



⑫

EUROPEAN PATENT APPLICATION

㉑ Application number: 88309413.8

㉓ Int. Cl.4: C 07 D 233/64

C 07 D 233/58, A 61 K 31/415

㉒ Date of filing: 07.10.88

㉓ Priority: 09.10.87 GB 8723715

㉔ Applicant: Farmos-Yhtymä Oy
P.O. Box 425
SF-20101 Turku 10 (FI)

㉔ Date of publication of application:
12.04.89 Bulletin 89/15

㉔ Inventor: Karjalainen, Arto Johannes
Myllyojantie 13 H 37
SF-90650 Oulu (FI)

㉔ Designated Contracting States:
AT BE CH DE ES FR GB GR IT LI LU NL SE

Kangas, Lauri Veikko Matti
Pasanteri 3
SF-21200 Raisio (FI)

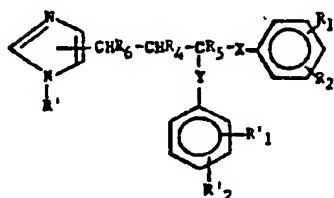
Kurkela, Kauko Oliva Antero
Pampinkuja 1C
SF-20900 Turku (FI)

㉔ Representative: Collier, Jeremy Austin Grey et al
J.A.Kemp & Co. 14, South Square Gray's Inn
London WC1R 5EU (GB)

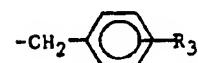
㉕ Aromatase Inhibiting 4(5)-Imidazoles.

R₈ is H or OH or one of R₅ and R₆ is H and the other, together with R₄, forms a bond and X and Y, which can be the same or different, are a bond, a straight C₁₋₂-alkyl or the corresponding alkenyl, and pharmaceutically acceptable salts thereof exhibit valuable pharmacological properties, especially aromatase inhibiting effects and are useful in the treatment of estrogen dependent diseases, e.g. breast cancer. Antimycotic and antifungal properties have also been found.

㉖ Imidazole derivatives of the formula:



wherein R₁, R₂, R'₁, and R'₂ which can be the same or different, are H, CH₃, C₂H₅, OCH₃, OH, CH₂OH, NH₂ or halogen; R'₁ is H or



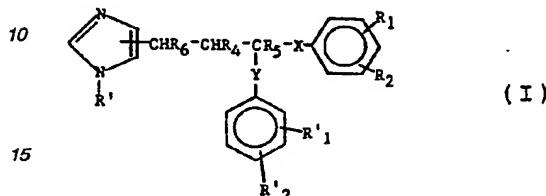
where R₃ is H, CH₃, or halogen; R₄ is H and R₅ is H or OH and

Description

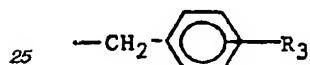
AROMATASE INHIBITING 4(5)-IMIDAZOLES

The present invention relates to substituted imidazole derivatives and their non-toxic, pharmaceutically acceptable acid addition salts, and their preparation, to pharmaceutical compositions containing the same and to their use.

5 The imidazole derivatives of the present invention have the general formula:



20 wherein R₁, R₂, R'₁ and R'₂, which can be the same or different, are H, CH₃, C₂H₅, OCH₃, OH, CH₂OH, NH₂ or halogen; R' is H or



30 where R₃ is H, CH₃ or halogen; R₄ is H and R₆ is H or OH and R₅ is H or OH or one of R₅ and R₆ is H and the other forms, together with R₄, a bond, and X and Y, which can be the same or different, are a bond, a straight C₁₋₂-alkyl or the corresponding alkenyl.

35 The non-toxic pharmaceutically acceptable acid addition salts of these compounds are also within the scope of the invention.

36 The compounds of the formula (I) form acid addition salts with both organic and inorganic acids. They can thus form many pharmaceutically usable acid addition salts, as, for instance, chlorides, bromides, sulfates, nitrates, phosphates, sulfonates, formates, tartrates, maleates, citrates, benzoates, salicylates, ascorbates and the like.

40 The invention includes within its scope pharmaceutical compositions comprising at least some of the compounds of formula (I) or a non-toxic, pharmaceutically acceptable salt thereof, and a compatible pharmaceutically acceptable carrier therefor.

45 The invention provides, for example, the following specific compounds of formula (I):

- 4-(3,3-diphenyl-3-hydroxypropyl)-1H-imidazole
- 4-[3,3-bis(4-chlorophenyl)-3-hydroxypropyl]-1H-imidazole
- 4-[3,3-bis(2-methylphenyl)-3-hydroxypropyl]-1H-imidazole
- 4-[3,3-bis(3-methylphenyl)-3-hydroxypropyl]-1H-imidazole
- 4-(3,3-diphenylpropen-2-yl)-1H-imidazole
- 4-[3,3-bis(4-chlorophenyl)propen-2-yl]-1H-imidazole
- 4-[3,3-bis(2-methylphenyl)propen-2-yl]-1H-imidazole
- 4-[3,3-bis(3-methylphenyl)propen-2-yl]-1H-imidazole
- 50 4-(3,3-diphenylpropyl)-1H-imidazole
- 4-[3,3-bis(2-methylphenyl)propyl]-1H-imidazole
- 1-benzyl-5-[3,3-bis(4-chlorophenyl)-3-hydroxypropyl]-1H-imidazole
- 1-benzyl-5-[5-(2,6-dimethylphenyl)-3-hydroxy-3-(2,6-dimethylphenylethyl)pentyl]-1H-imidazole
- 1-benzyl-5-[3,3-bis(4-chlorophenyl)propen-2-yl]-1H-imidazole
- 55 4-[3-(4-chlorophenyl)-3-hydroxy-3-phenylpropyl]-1H-imidazole
- 1-benzyl-4-(3,3-diphenylpropyl)-1H-imidazole
- 1-benzyl-5-(3,3-diphenylpropyl)-1H-imidazole
- 4-[5-(2,6-dimethylphenyl)-3-(2,6-dimethylphenylethyl)pentyl]-1H-imidazole
- 4-[3,3-bis(3-methylphenyl)propyl]-1H-imidazole
- 60 1-(4-chlorobenzyl)-4-(3,3-diphenylpropyl)-1H-imidazole
- 1-(4-chlorobenzyl)-5-(3,3-diphenylpropyl)-1H-imidazole
- 4-[5-(2,6-dimethylphenyl)-3-hydroxy-3-(2,6-dimethylphenylethyl)pentyl]-1H-imidazole
- 4-[3,3-bis(3-fluorophenyl)propen-2-yl]-1H-imidazole

4-[3,3-bis(3-fluorophenyl)propyl]-1H-imidazole
 4-[3,3-bis(3,5-dimethylphenyl)propyl]-1H-imidazole
 1-benzyl-5-(3,3-diphenylpropen-2-yl)-1H-imidazole
 1-benzyl-5-[3,3-bis(3,5-dimethylphenyl)-3-hydroxypropyl]-1H-imidazole
 1-benzyl-5-[3,3-bis(3,5-dimethylphenyl)propen-2-yl]-1H-imidazole
 1-benzyl-5-[3,3-bis(2-methoxyphenyl)propen-2-yl]-1H-imidazole
 1-benzyl-5-[3,3-bis(3-methoxyphenyl)propen-2-yl]-1H-imidazole
 1-benzyl-5-[3,3-bis(4-methoxyphenyl)propen-2-yl]-1H-imidazole
 1-benzyl-5-[3,3-bis(2,3-dimethylphenyl)propen-2-yl]-1H-imidazole
 1-benzyl-5-[3,3-bis(2-methylphenyl)propen-2-yl]-1H-imidazole
 1-benzyl-5-[3,3-bis(4-methylphenyl)propen-2-yl]-1H-imidazole
 1-benzyl-5-[3,3-bis(3,5-dimethylphenyl)propyl]-1H-imidazole
 1-benzyl-5-[3,3-bis(3-methoxyphenyl)propyl]-1H-imidazole
 4-[3,3-bis(3,5-dimethylphenyl)propyl]-1H-imidazole
 4-[3,3-bis(2,3-dimethylphenyl)propyl]-1H-imidazole
 4-[3,3-bis(2-methoxyphenyl)propyl]-1H-imidazole
 4-[3,3-bis(4-methoxyphenyl)propyl]-1H-imidazole
 4-[3,3-bis(4-methylphenyl)propyl]-1H-imidazole

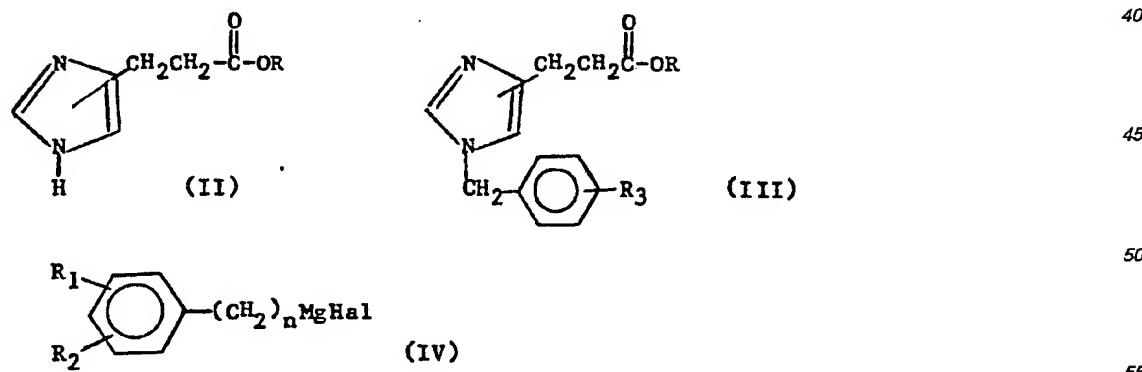
5
10
15
20
25

The compounds of the present invention have been found to possess aromatase inhibiting properties and are therefore valuable in the treatment of estrogen dependent diseases, e.g. breast cancer. Antimycotic and antifungal properties have also been found.

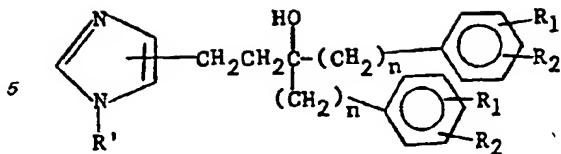
According to a feature of the Invention, the compounds of formula I wherein the branches



are identical are prepared by a successive sequence of reactions comprising a Grignard reaction of 4(5)-imidazole propionic acid alkyl ester (II) or its 1-benzyl derivative III with an appropriate aryl- or arylalkylmagnesium halide IV following the loss of water and hydrogenation



In the formulae (II) to (IV) R is alkyl, R₃ is H, CH₃ or halogen, n is 0 to 2 and R₁ and R₂, which can be the same or different, are H, CH₃, C₂H₅, OCH₃, OH, CH₂OH, NH₂ or Hal (Hal = halogen). The first reaction step, the Grignard-reaction, leads to the following compounds of formula (I):



10

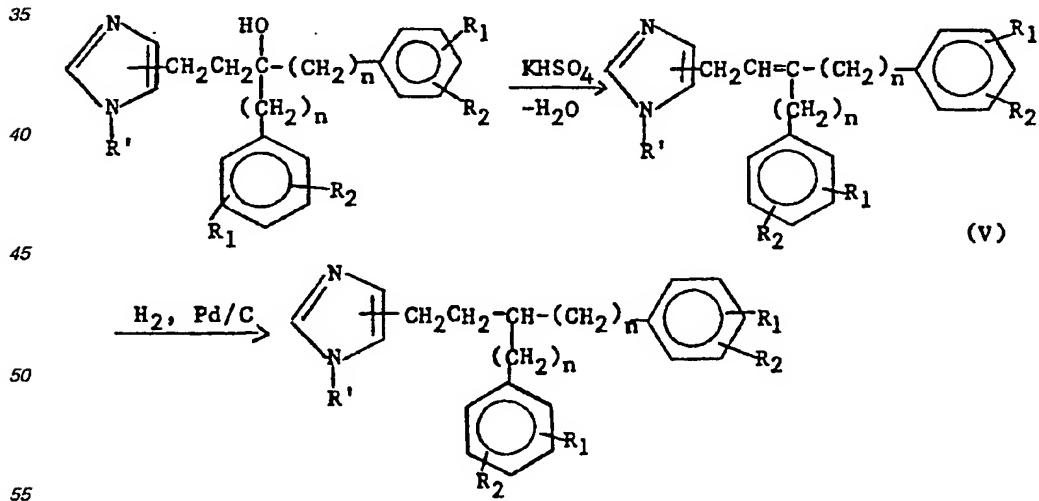
In this reaction the arylalkylmagnesium halide derivative can be, for example, an arylalkylmagnesiumbromide derivative, which is prepared by reacting the corresponding arylalkylbromide derivative with magnesium. Suitable solvents for the reaction include a variety of ethers, preferably tetrahydrofuran.

15 The arylalkylmagnesiumhalide derivative is prepared in the usual way by adding the arylalkylhalide derivative in a suitable solvent, e.g. tetrahydrofuran, dropwise onto magnesium turnings covered by tetrahydrofuran, at the boiling point of the reaction mixture. When the magnesium turnings have reacted, the mixture is cooled slightly and the 4(5)-imidazole propionic acid alkyl ester or its 1-benzylsubstituted derivative is added in solid form in small portions or dropwise in tetrahydrofuran.

20 After the addition, the reaction mixture is refluxed until all of the 4(5)-imidazole derivative has reacted. The reaction time varies between one and five hours.

Further according to the feature of the invention, the compounds of formula (I), wherein R₄ and R₅ both are hydrogen or together form a bond, are prepared by dehydration of the compounds of formula (I), where R₅ is OH, and by catalytic addition of hydrogen in the second step. Water is eliminated by usual methods, i.e. by heating with concentrated hydrochloric acid or by heating with dry potassium hydrogen sulfate. The unsaturated derivatives (V) (the compounds of formula (I) wherein R₄ and R₅ together form a bond) are isolated and after that hydrogenated. Alternatively they can be hydrogenated directly in an acid medium without previous isolation. The hydrogenation is conveniently carried out at room temperature with good stirring in alcohol, e.g. ethanol in the presence of a catalyst in a hydrogen atmosphere. Suitable catalysts are for example platinum oxide, palladium-on-carbon or Raney-nickel.

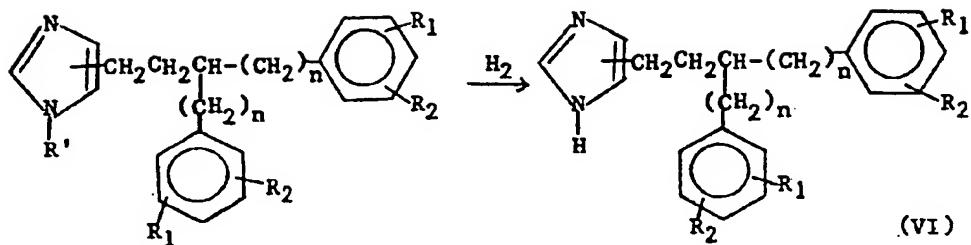
30 The reaction scheme for these steps can be illustrated as follows:



60 If R' is a substituted or unsubstituted benzyl, this group may be removed by hydrogenation as well. In this case the hydrogenation is performed in an acidic medium such as hydrochloric acid-ethanol mixture at elevated temperature.

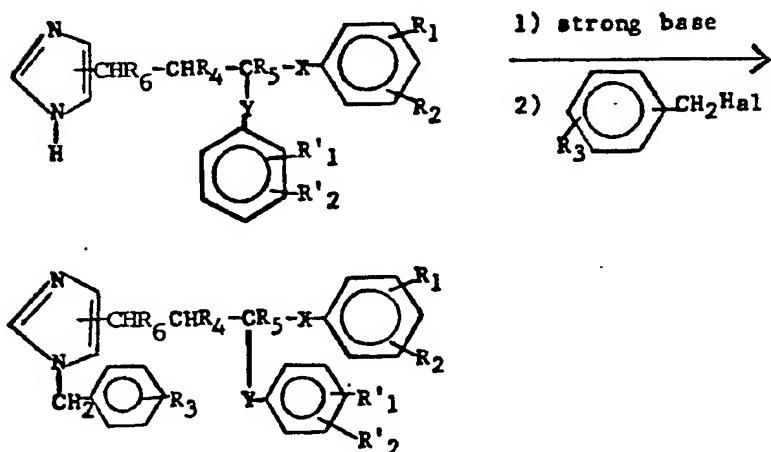
The reaction scheme of this hydrogenation which leads to compounds of formula (I) wherein R₄ and R₅ both are hydrogen can be illustrated as follows:

65

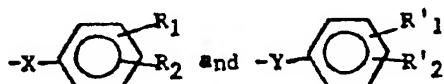


The compound (VI) can also be prepared directly from the compounds (V) by hydrogenating both the double bond and the protecting benzyl group at the same time.

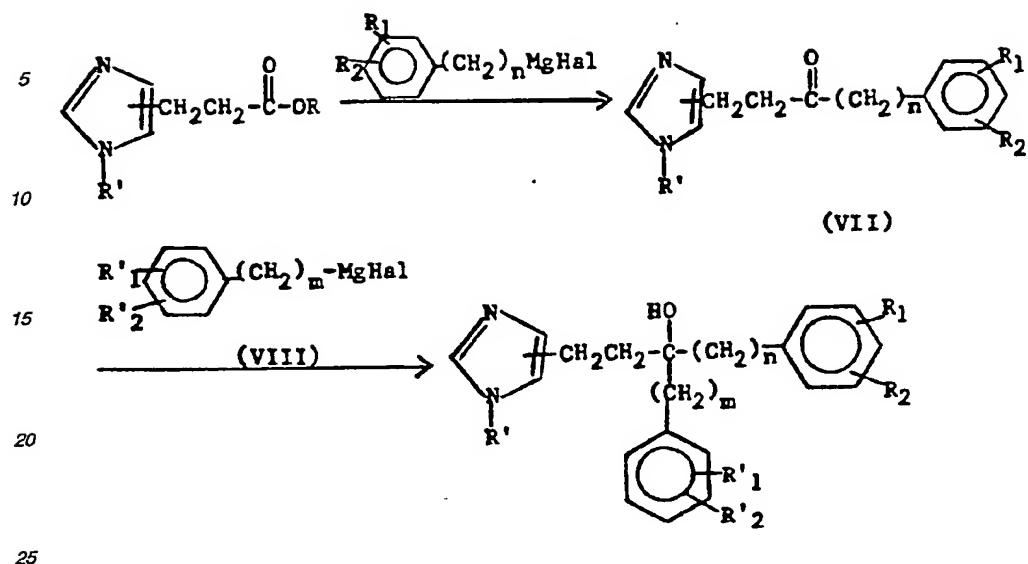
Another method for the preparation of compounds of formula (I) where R' is a benzyl is the benzylation of the corresponding compound where R' is hydrogen. The starting compound is first treated with a strong base such as sodium hydroxide in water or sodium hydride in an appropriate solvent, e.g. dimethyl formamide to give the alkali metal salt of the imidazole and then in the second step adding to this benzyl halide. The reaction scheme can be illustrated as follows:



Yet another process for the preparations of compounds of formula (I) wherein the branches



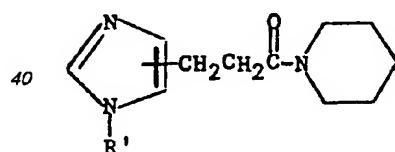
are different, comprises in the first stage a series of two successive Grignard reactions starting from 4(5)-imidazolylpropionic acid alkyl ester or from 1-benzyl-4(5)-imidazolylpropionic acid alkyl ester as previously. Now, however, the amount of the Grignard reagent is reduced as well as the reaction temperature, to stop the reaction at the ketone stage to give the 4(5)-imidazolylpropyl aryl or arylalkyl ketone (VII), which further is reacted with another Grignard reagent (VIII) to give a compound of the formula (I) where R5 is OH. The reactions are illustrated as follows:



In the reaction scheme above m and n, which can be the same or different, are 0 to 2.

Choosing appropriate conditions for the dehydration of the compounds of formula (I) where R₅ is OH results 30 in the corresponding compounds of formula (I) where one of the alkyl chains X or Y is transformed to the corresponding alkanyl chain.

In order to achieve a better control of the reactions above, as starting material may be used an amide of the 4(5)-imidazole propionic acid as well. Especially suitable in this respect is for example a piperidinyl amide of the formula 35



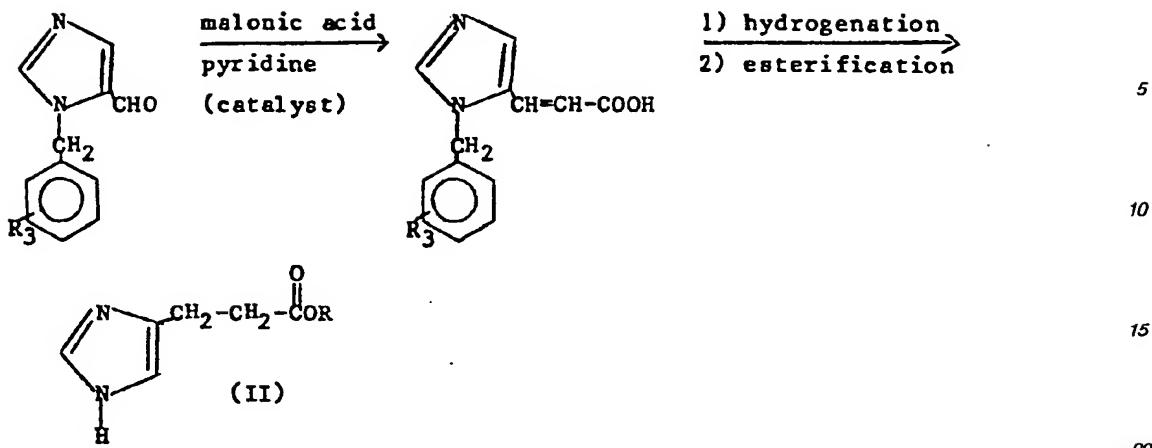
45

In the processes described above, the 4(5)-imidazole propionic acid esters (II) and (III) may be prepared for example starting from 1-benzyl-5-imidazole carbaldehyde and malonic acid, which are condensed together to 50 form a 5-(1-benzylimidazole)acrylic acid. When this compound is hydrogenated under acidic conditions at elevated temperature (70-80°C) in the presence of a catalyst, 4-imidazole propionic acid is formed. The subsequent treatment with alcohol, e.g. methanol, in the presence of dry hydrochloric acid leads to 4-imidazole propionic acid alkyl ester, which is used as starting material in the Grignard reaction:

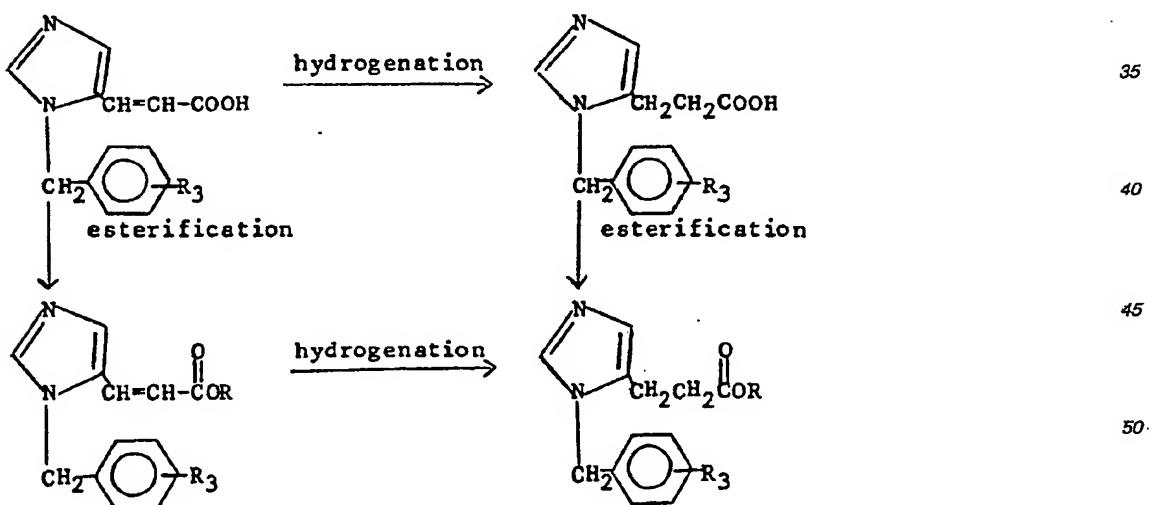
55

60

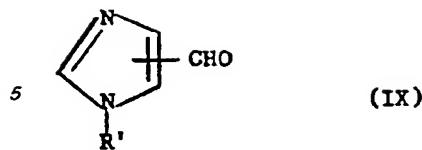
65



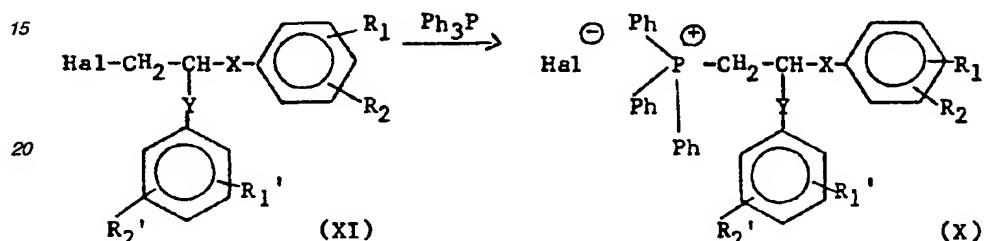
When 5-(1-benzylimidazole) acrylic acid (the benzyl group may be substituted or unsubstituted) is hydrogenated at room temperature in alcohol 1-benzyl-5-imidazole propanoic acid is achieved. The following treatment with alcohol in the presence of dry hydrochloric acid at elevated temperature leads to another possible starting material for the Grignard reaction, namely 1-benzyl-5-imidazole propionic acid alkyl ester. The described reaction steps can be conducted in the opposite order as well. The reaction schemes are as follows:



The compounds of formula (I) can be prepared by the Wittig reaction and the Grignard reaction wherein the starting compound is a 4(5)-imidazole aldehyde (IX). In the formula (IX) R' is as defined before.



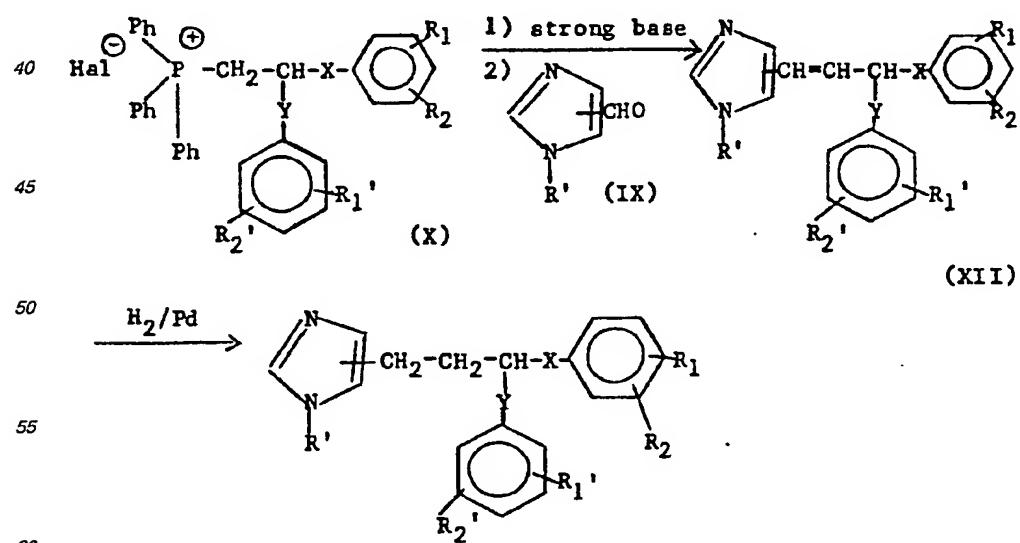
10 In the Wittig reaction the first step is to prepare a phosphonium salt (X) from the corresponding halogenated hydrocarbon (XI) by reacting it with triphenylphosphine. The reaction scheme can be illustrated as follows:



in which R', R', R₁, and R₂ are as hereinbefore defined.

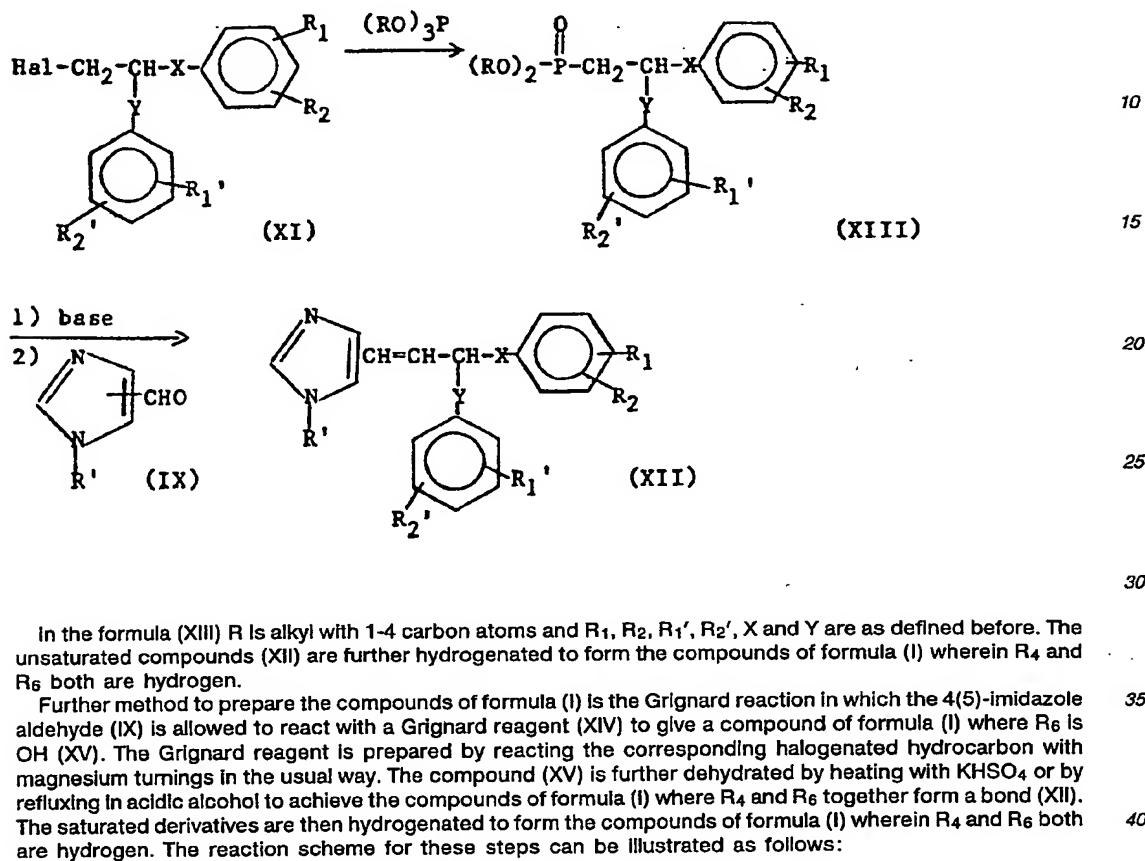
30 In the second step of the Wittig reaction the compound (X) is treated with a strong base to form a phosphorus ylide which is further allowed to react with the 4(5)-imidazole aldehyde (IX) to achieve the compounds of formula (I) wherein R₄ and R₆ together form a bond (XII). The strong base can be NaH or BuLi in a proper solvent such as dimethoxyethane, tetrahydrofuran or DMF. Further alkali metal alkoxides the corresponding alcohols as solvent and NaH in DMSO can be used as proton acceptors. the compounds (XII) are isolated and after that hydrogenated as has been described before to achieve the compounds of formula (I) wherein R₄ and R₆ both are hydrogen. The reaction scheme for these steps can be illustrated as follows:

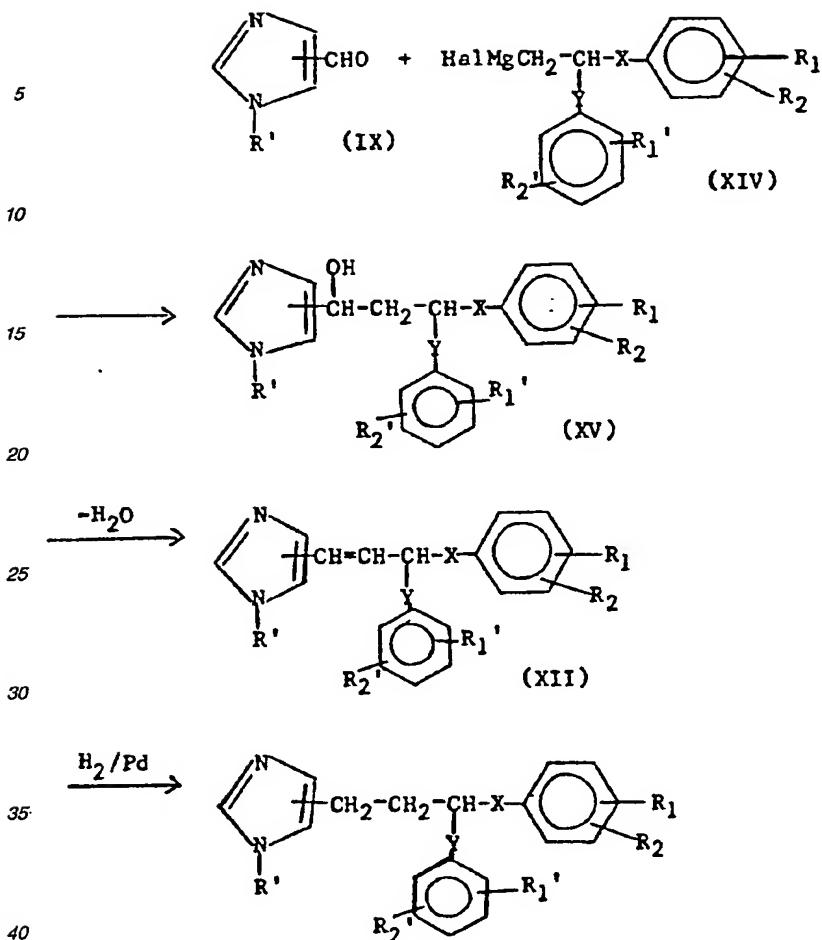
35



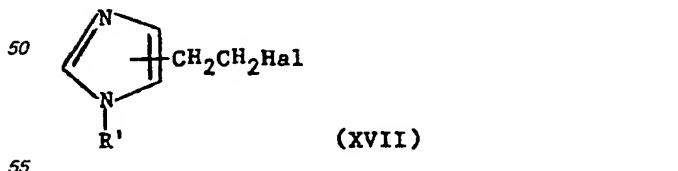
65 The compounds of formula (I) can also be prepared by a modified Wittig reaction, namely the Horner-Emmons or Wadsworth-Emmons reaction where the phosphonate (XIII) which is prepared from the halogenated hydrocarbon (XI) and a triester of phosphonic acid (e.g. (EtO)₃P) by the Arbuzow reaction reacts

firstly with a base (e.g. NaH in DMSO or in dimethoxyethane) and then with the aldehyde (IX). The product (XII) formed is a compound of formula (I) where R₄ and R₆ together form a bond. The reaction scheme can be illustrated as follows:





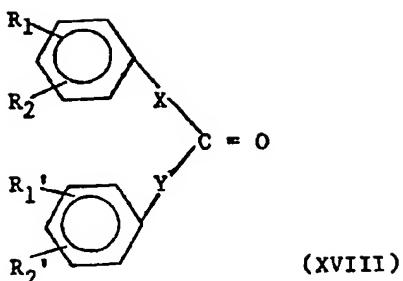
45 Further the compounds of formula (I) can be prepared by a Grignard reaction where the Grignard reagent (XVI) is prepared from a 4(5)-imidazolylalkylhalogenide (XVII)



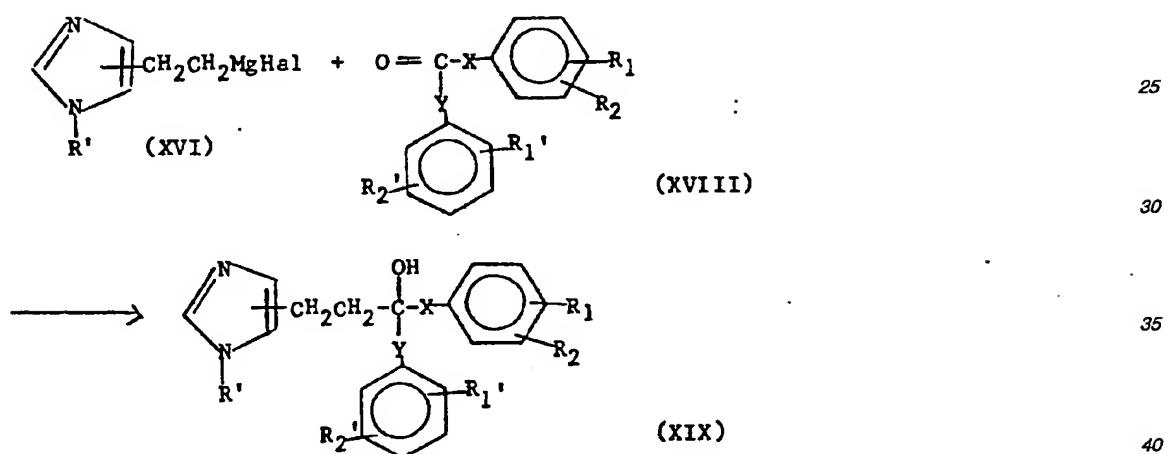
by allowing it to react firstly with magnesium and then with a suitable ketone (XVIII)

60

65



The reaction scheme of this reaction which leads to compounds of formula (I) where R_5 is OH (XIX) can be illustrated as follows:



The compounds (XIX) can further be dehydrated and hydrogenated as described before to achieve the compounds of formula (I) wherein R_4 and R_5 both are hydrogen.

Administration of compounds of formula (I), their non-toxic, pharmaceutically acceptable acid salts or mixtures thereof may be achieved parenterally, intravenously or orally. Typically, an effective amount of the derivative is combined with a suitable pharmaceutical carrier. As used herein, the term "effective amount" encompasses those amounts which yield the desired activity without causing adverse side-effects. The precise amount employed in a particular situation is dependent upon numerous factors such as method of administration, type of mammal, condition for which the derivative is administered, etc., and of course the structure of the derivative.

The pharmaceutical carriers which are typically employed with the derivatives of the present invention may be solid or liquid and are generally selected with the planned manner of administration in mind. Thus, for example, solid carriers include lactose, sucrose, gelatin and agar, while liquid carriers include water, syrup, peanut oil and olive oil. Other suitable carriers are well-known to those skilled in the art of pharmaceutical formulations. The combination of the derivative and the carrier may be fashioned into numerous acceptable forms, such as tablets, capsules, suppositories, solutions, emulsions, and powders.

The compounds of the invention are especially valuable as aromatase inhibiting agents and are therefore useful in the treatment of estrogen dependent diseases, e.g. breast cancer.

Estrogens are essential steroids in the physiology and function of normal development of breast and sex organs in women. On the other hand estrogens are known to stimulate the growth of estrogen dependent cancers, especially breast and endometrial cancers, and they may increase the risk of development of breast cancer if given at pharmacological doses for a long time.

Excessive production of estradiol may also cause other, benign disorders in hormone dependent organs.

The importance of estrogens as cancer growth stimulators and/or regulators is clearly stressed by the fact that antiestrogens have reached a central position in the treatment of estrogen receptor rich breast cancers. Antiestrogens act by binding to estrogen receptors and thereby inhibiting the biological effects of estrogens. Another approach for blocking estrogen effect is to inhibit the synthesis of estrogens. This has been achieved 5 clinically by the unspecific steroid synthesis inhibitor aminoglutethimide. The estrogen synthesis could be blocked specifically by inhibiting the enzyme aromatase, which is the key enzyme in biochemical estrogen synthesis pathway. Aromatase inhibition seems highly promising because several breast tumours synthesise estradiol and estrone in situ and exhibit therefore continuous growth stimulation (Alan Lipton et al., Cancer 59: 779-782, 1987).

10 The ability of the compounds of the invention to inhibit the enzyme aromatase was shown by the in vitro assay method according to M. Pasanen, Biological Research in Pregnancy, vol. 6, No. 2, 1985 (pp. 94-99). Human aromatase enzyme was used. The enzyme was prepared from human placenta, which is rich in the enzyme. Microsomal fraction (100000 x g precipitate) was prepared by centrifugation. The enzyme preparation was used without further purification. Test compounds listed in Table 1 were added with 100000 dpm of 15 1,2-[³H]-androstene-3,17-dione and NADPH generating system. The concentrations of the test compounds were 0,001; 0,01; 0,1 and 1,0 mM. The incubation was carried out at 37°C for 40 min. Aromatization of 1,2-[³H]-androstene-3,17-dione results in the production of ³H₂O. The tritiated water and the tritiated substrate are easily separated by a Sep-Pak® minicolumn, which absorbs the steroid but allows free water elution. 20 Radioactivity was counted by a liquid scintillation counter. Aromatase inhibition was evaluated by comparing the ³H₂O-radioactivity of inhibitor treated samples to controls containing no inhibitor. IC-10, IC-50 and IC-90 values were calculated as concentrations which inhibited the enzyme activity 10%, 50% and 90%, respectively. These concentrations are presented in Table 2.

Table 1

25 Compounds tested

No.	Name
30	1. 4-[5-(2,6-dimethylphenyl)-3-hydroxy-3-(2,6-dimethylphenylethyl)-pentyl]-1H-imidazole
	2. 4-[3,3-bis(4-chlorophenyl)-3-hydroxypropyl]-1H-imidazole
	3. 4-(3,3-diphenyl-3-hydroxypropyl)-1H-imidazole
	4. 4-(3,3-diphenylpropen-2-yl)-1H-imidazole
	5. 4-(3,3-diphenylpropyl)-1H-imidazole
35	6. 4-[3,3-bis(2-methylphenyl)-3-hydroxypropyl]-1H-imidazole
	7. 4-[3,3-bis(4-chlorophenyl)propen-2-yl]-1H-imidazole
	8. 4-[3,3-bis(2-methylphenyl)propen-2-yl]-1H-imidazole
	9. 4-[3,3-bis(2-methylphenyl)propyl]-1H-imidazole
	10. 1-benzyl-5-(3,3-diphenylpropyl)-1H-imidazole
40	11. 4-[3,3-bis(3-methylphenyl)propyl]-1H-imidazole
	12. 4-[3,3-bis(3-methylphenyl)propen-2-yl]-1H-imidazole
	13. 4-[3,3-bis(3-methoxyphenyl)propyl]-1H-imidazole
	14. 4-[3,3-bis(2,3-dimethylphenyl)propyl]-1H-imidazole
45	15. 1-benzyl-5-[3,3-bis(3-methoxyphenyl)propyl]-1H-imidazole
	16. 1-benzyl-5-[3,3-bis(3-methoxyphenyl)propen-2-yl]-1H-imidazole
	17. 4-[3,3-bis(3,5-dimethylphenyl)propyl]-1H-imidazole
	18. 4-[3,3-bis(4-methylphenyl)propyl]-1H-imidazole
	19. 4-[3,3-bis(3-fluorophenyl)propyl]-1H-imidazole
50	20. 1-benzyl-5-(3,3-diphenylpropen-2-yl)-1H-imidazole

55

60

65

Table 2

Inhibition of human aromatase by test compounds. IC-10, IC-50 and IC-90 represent the concentration which inhibit the enzyme by 10%, 50 % and 90 %, respectively

Compound No.	IC-10 mmol/l	IC-50 mmol/l	IC-90 mmol/l	
1	0,02	1,0	> 1	10
2	0,004	0,06	1,0	
3	0,004	0,07	1,0	
4	< 0,001	0,006	0,10	
5	0,0015	0,015	0,40	
6	0,015	0,30	> 1	15
7	0,002	0,030	0,6	
8	0,0015	0,080	> 1	
9	0,002	0,030	0,6	
10	~ 0,0006	0,004	0,10	20
11	~ 0,0006	0,006	0,10	
12	0,001	0,080	1	
13		0,003		
14		0,044		
15		0,004		25
16		0,014		
17		0,130		
18		0,014		
19		0,019		30
20		0,050		

The daily dose for a patient varies from about 20 to about 200 mg, administered orally.

The toxicity of the imidazole derivatives of the present invention was studied in rats. There were 5 female rats in each drug group and dosing was carried out during 8 days. The dose level used was 10 mg/kg/day orally. The derivatives tested were 4-(3,3-diphenylpropen-2-yl)-1H-imidazole, 4-(3,3-diphenylpropyl)-1H-imidazole, 4-[3,3-bis(2-methylphenyl)propyl]-1H-imidazole, 1-benzyl-5-(3,3-diphenylpropyl)-1H-imidazole and 1-benzyl-5-(3,3-diphenylpropen-2-yl)-1H-imidazole.

The behaviour, appearance and mortality of the animals were followed daily. The animals were weighed before and after dosing period. The organs were examined macroscopically at autopsy. The liver, uterus and ovaries were weighed. No mortality were observed. The weight development was normal in all groups. In the groups that were treated with 4-(3,3-diphenylpropen-2-yl)-1H-imidazole and 1-benzyl-5-(3,3-diphenylpropen-2-yl)-1H-imidazole a slight piloerection was observed probably as pharmacological effects of the drugs. No drug-related findings were observed in organ weights or in macroscopical pathology. In conclusion, all studied compounds were well tolerated.

The following examples illustrate the invention.

Example 1

4-(3,3-diphenyl-3-hydroxypropyl)-1H-imidazole

a) 5-(1-benzylimidazole)acrylic acid

In a flask are placed 18,6 g of 5-(1-benzylimidazole)-carbaldehyde, 10,4 g of malonic acid, and 4,8 ml of pyridine. The mixture is heated on a boiling water bath for 16 hours. It is then cooled and diluted with water. The precipitate which is the product is filtered and washed with water. Yield 15 g. M.p.221-226°C. ⁵⁵
¹H NMR: 5.15 (s, 1H), 5.64 (s, 2H), 6.58 (d, 1H), 7.3-7.5 (m, 5H), 7.61 (d, 1H), 8.08 (s, 1H), 9.07 (s, 1H)

b) 4(5)-imidazole propionic acid ethyl ester

5-(1-benzylimidazole)acrylic acid (15 g) is dissolved in 50 ml of 4-N hydrochloric acid. About 60 mg of 10 % Pd/C are added and the mixture is stirred vigorously under a hydrogen atmosphere at about 85°C until no more hydrogen is consumed. The reaction mixture is then filtered and evaporated to dryness.

The residue is dissolved in 50 ml of abs. ethanol and dry hydrogen chloride gas is passed into the solution for 4 hours during which time the reaction mixture is maintained at reflux with stirring. The mixture is then evaporated to dryness to give an oily residue which is a crude product useful as such in the following Grignard reaction.

¹H NMR: 1.237 (t, 3H), 2.656 (t, 2H), 2.936 (t, 2H), 4.137 (q, 2H), 6.804 (s, 1H), 7.559 (s, 1H)

c) 4-(3,3-diphenyl-3-hydroxypropyl)-1H-imidazole

3,3 g of magnesium turnings are covered with 100 ml of dry tetrahydrofuran. To that mixture is then added dropwise a solution of 21,8 g of bromobenzene in 30 ml of dry tetrahydrofuran at such a rate that a smooth reaction is maintained. After the addition is complete, the reaction mixture is refluxed for one additional hour and cooled to room temperature. The reaction mixture is then added dropwise to a solution of 4(5)-imidazole propionic acid ethyl ester (7,8 g) in 50 ml of tetrahydrofuran at room temperature. After the addition is complete, the reaction mixture is stirred for an additional hour at 40-50°C. The mixture is then cooled and poured into cold water. Tetrahydrofuran is evaporated and to the solution is added conc. hydrochloric acid (20 ml). The solution is cooled and the precipitate which contains the product as hydrochloride salt is removed by filtration, washed and dried. Yield 11,2 g. M.p. 189-191°C.

¹H NMR: 2.703 (s, 4H), 4.758 (s, 3H), 7.214-7.429 (m, 11H), 8.457 (s, 1H)

In the same way, via the Grignard reaction starting from 4(5)-imidazole propionic acid ethyl ester and from proper substituted bromobenzene, can also be prepared other compounds of the invention.

For example the following substituted derivatives were prepared:

4-[3,3-bis(4-chlorophenyl)-3-hydroxypropyl]-1H-imidazole. M.p. of hydrochloride 85-89°C.

4-[3,3-bis(2-methylphenyl)-3-hydroxypropyl]-1H-imidazole. M.p. of hydrochloride 211-213°C.

4-[3,3-bis(3-methylphenyl)-3-hydroxypropyl]-1H-imidazole. M.p. of hydrochloride 170-172°C.

20

Example 2

4-(3,3-diphenylpropen-2-yl)-1H-imidazole

2,0 g of 4-(3,3-diphenyl-3-hydroxypropyl)-1H-imidazole hydrochloride is mixed with 20 g of anhydrous potassium hydrogen sulfate and the mixture is warmed on an oil bath at 150-155°C for 4 hours. The mixture is then cooled and 20 ml water is added. The mixture is made alkaline with sodium hydroxide solution and cooled. The precipitate, which is the product, is filtered, washed with water and dried. Yield 1,25 g. After recrystallization from water-ethanol, the product melts at 124-128°C.

¹H NMR: 3.42 (d, 2H), 4.756 (s, 1H), 6.284 (t, 1H), 6.768 (s, 1H), 7.2-7.4 (m, 10H), 7.559 (s, 1H)

30 According to the same procedure for example the following substituted derivatives were prepared:

4-[3,3-bis(4-chlorophenyl)propen-2-yl]-1H-imidazole hydrochloride. M.p. 158-163°C.

4-[3,3-bis(2-methylphenyl)propen-2-yl]-1H-imidazole hydrochloride. M.p. 195-198°C.

4-[3,3-bis(3-methylphenyl)propen-2-yl]-1H-imidazole. M.p. 115-118°C.

4-[3,3-bis(3-fluorophenyl)propen-2-yl]-1H-imidazole. M.p. of hydrochloride is 125-128°C.

35

Example 3

4-(3,3-diphenylpropyl)-1H-imidazole

40 4-(3,3-diphenylpropen-2-yl)-1H-imidazole (0,7 g) is dissolved in ethanol and a catalytic amount of Pd/C (10 %) is added. The reaction mixture is agitated vigorously at room temperature in a hydrogen atmosphere until the uptake of hydrogen ceases. The mixture is filtered and the filtrate is evaporated to dryness. The residue is recrystallized from water-ethanol mixture. Yield 0,4 g, m.p. 115-117°C.

¹H NMR: 2.3-2.5 (m, 4H), 3.919 (t, 1H), 4.752 (s, 1H), 6.708 (s, 1H), 7.1-7.3 (m, 10H), 7.532 (s, 1H)

According to the same procedure as the example the following substituted derivatives were prepared:

4-[3,3-bis(4-chlorophenyl)propyl]-1H-imidazole, hydrochloride. M.p. 84-87°C.

4-[3,3-bis(2-methylphenyl)propyl]-1H-imidazole. M.p. 111-114°C.

1H NMR (as base):

2.272 (s, 6H), 2.2-2.5 (m, 4H), 3.823 (t, 1H), 6.691 (s, 1H), 6.8-7.2 (m, 8H), 7.440 (s, 1H).

4-[3,3-bis(3-fluorophenyl)propyl]-1H-imidazole

50 1H NMR (as HCl):

2.3-2.8 (m, 4H), 4.060 (t, 1H), 4.784 (s, 2H), 6.7-7.4 (m, 9H), 8.743 (s, 1H)

Example 4

55 1-benzyl-5-[3,3-bis(4-chlorophenyl)-3-hydroxypropyl]-1H-imidazole

a) 1-benzyl-5-imidazole acrylic methyl ester

60 In a flask are placed 12,0 g of 5-(1-benzylimidazole)acrylic acid (prepared in example 1), 70 ml of methanol and dry hydrogen chloride gas is passed into the solution for 4 hours, during which time the reaction mixture is maintained at reflux. The mixture is then evaporated to dryness and the residue is dissolved in cold water. The solution is then made alkaline with sodium carbonate and the precipitate, which is the product, is filtered, washed with water and dried. Yield 12,2 g; m.p. 137-139°C.

¹H NMR: 3.781 (s, 3H), 5.490 (s, 2H), 6.462 (d, 1H), 7.2-7.5 (m, 5H), 7.493 (d, 1H), 7.710 (s, 1H), 8.083 (s, 1H)

b) 1-benzyl-5-imidazole propionic acid methyl ester

The double bond of the side chain is hydrogenated in abs. ethanol Pd/C as catalyst. When the uptake of hydrogen ceases, the reaction mixture is filtered and the filtrate is evaporated to dryness. The residue is dissolved in methylene chloride, which is washed with water. Methylene chloride phase is then dried and evaporated to dryness to give the product, which is used as such in the accompanying Grignard reactions. ¹³C NMR: Aliphatic carbons are detected at ppm: 19.374, 32.573, 48.466, 51.675; aromatic carbons are detected at ppm: 126.569, 128.022, 128.748, 128.960, 130.474, 136.074, 137.88; and carbonyl at ppm: 172/522

5

c) 1-benzyl-5-[3,3-bis(4-chlorophenyl)-3-hydroxypropyl]-1H-imidazole

The Grignard reagent is prepared from 2,4 g of magnesium turnings and from 19,1 g of p-chlorobromobenzene in tetrahydrofuran as is described in Example 1 c).

10

1-benzyl-5-imidazole propionic acid methyl ester (6,4 g) in tetrahydrofuran is heated at 60°C and to this is then added dropwise p-chlorophenylmagnesium bromide prepared above. After the addition is complete, the reaction mixture is refluxed for an additional 3 hours, cooled and poured into cold water. Tetrahydrofuran is evaporated, toluene is added and the mixture is made acidic with hydrochloric acid. The precipitated product is filtered, washed with ether and dried. Yield 12,2 g; m.p. 210-213°C. M.p. of nitrate 157-160°C (made in water-ether mixture). M.p. of hydrochloride (from ethylacetate) 178-187°C.

15

¹H-NMR: 2.985 (s, 4H), 4.854 (s, 2H), 5.330 (s, 2H), 7.06-7.46 (m, 14H), 8.993 (s, 1H)

Other 1-benzyl substituted derivatives are also prepared in the same way. For example: 1-benzyl-5-[5-(2,6-dimethylphenyl)-3-hydroxy-3-(2,6-dimethylphenylethyl)pentyl]-1H-imidazole from 1-benzyl-5-imidazole propionic acid methyl ester and 2-(2,6-dimethylphenyl)ethylmagnesium bromide. Melting point of the hydrochloride is 67-71°C.

20

Example 5

25

1-benzyl-5-[3,3-bis(4-chlorophenyl)propen-2-yl]-1H-imidazole

4,1 g of 1-benzyl-[3,3-bis(4-chlorophenyl)-3-hydroxypropyl]-1H-imidazole and 22,0 g of anhydrous potassium hydrogen sulfate are heated at 150°C for 4 hours. The mixture is cooled, 100 ml of ethanol is added to dissolve the product. The mixture is then filtered and the filtrate is evaporated to minor volume. Water is added and the mixture is made basic with sodium hydroxide. The precipitate, which is the product, is filtered, washed with water and dried. The product is recrystallized from water-ethanol. Yield 2,3 g. Nitrate is made in water with nitric acid.

30

¹H NMR: 3,293 (d, 2H), 5.287 (s, 1H), 6.010 (t, 1H), 6.9-7.4 (m, 14H), 9.330 (s, 1H)

35

Example 6

4-[3-(4-chlorophenyl)-3-hydroxy-3-phenylpropyl]-1H-imidazole

a) 3-(4-imidazolyl)propyl 4-chlorophenyl ketone

40

0,85 g of magnesium turnings are covered with 20 ml of dry tetrahydrofuran, the mixture is heated to boiling and to it is added 6,8 g of 4-bromochlorobenzene in tetrahydrofuran at such a rate that a smooth reaction is maintained. After the addition is complete, the reaction mixture is refluxed for one additional hour. The reaction mixture is then cooled and added dropwise at room temperature to a solution of 4(5)-imidazole propionic acid ethyl ester (4,0 g) in tetrahydrofuran. After addition the reaction mixture is stirred for an additional hour at room temperature. It is then poured into cold water and made acidic with hydrochloric acid. The reaction mixture is then washed with methylene chloride, made alkaline with sodium hydroxide, and the product is extracted to methylene chloride. Yield 2,2 g. Hydrochloride salt is made in conc. hydrochloric acid. M.p. 160-161°C.

45

b) 4-[3-(4-chlorophenyl)-3-hydroxy-3-phenylpropyl]-1H-imidazole

50

Phenylmagnesiumbromide is made in tetrahydrofuran from 0,51 g of magnesium turnings and 3,3 g of bromobenzene. 3-(4-imidazolyl)propyl 4-chlorophenyl ketone (2,3 g) is dissolved in tetrahydrofuran and phenylmagnesiumbromide is dropped to that solution at room temperature. After addition the reaction mixture is stirred at 40-50°C for additional 3 hours. It is then cooled and poured into cold water. Water is made acidic with hydrochloric acid. The product is extracted into methylenechloride, which is evaporated into dryness. The product as hydrochloride is recrystallized from water-ethanol. Yield 3,2 g.

55

Example 7

1-benzyl-4-(3,3-diphenylpropyl)-1H-imidazole and 1-benzyl-5-(3,3-diphenylpropyl)-1H-imidazole

60

4-(3,3-diphenylpropyl)-1H-imidazole (2,6 g) is dissolved in 6 ml of dry dimethylformamide. While stirring 0,5 g of NaH (60%) is added during half an hour at room temperature. After addition the reaction mixture is stirred additional one hour. 1,7 g of benzylbromide in 3 ml of dimethylformamide is then dropped at room temperature and stirring is continued for 4 hours. The reaction mixture is poured to cold water (30 ml) and the mixture is extracted with toluene. Toluene extracts are then washed with water and evaporated to dryness. The residue, which is the mixture of products, is purified and separated to pure isomers by column chromatography

65

(methylene chloride/methanol, 9,5/0,5).

¹H NMR of the products:

One of the isomers:

2.57 (m, 4H), 3.52 (1H), 3,877 (t, 1H), 5.362 (s, 2H), 6.531 (s, 1H), 7.05-7.40 (m, 15H), 9.567 (s, 1H)

5 The other isomer:

2.375 (m, 4H), 3,858 (t, 1H), 5.253 (s, 2H), 7.01-7.36 (m, 16H), 9.441 (s, 1H)

Example 8

10 1-(4-chlorobenzyl)-4-(3,3-diphenylpropyl)-1H-imidazole and

1-(4-chlorobenzyl)-5-(3,3-diphenylpropyl)-1H-imidazole

The compounds were prepared in the same way as the compounds in Example 7 starting from 4-(3,3-diphenylpropyl)-1H-imidazole and 4-chlorobenzylchloride.

¹H NMR of the products:

15 One isomer:

2.48 (m, 4H), 3.934 (t, 1H), 4.999 (s, 2H), 6.514 (s, 1H), 7.0-7.3 (m, 14H), 7.517 (s, 1H)

The other isomer:

2.33 (m, 4H), 3.887 (t, 1H), 4.852 (s, 2H), 6.7-7.5 (m, 16H)

20 Example 9

4-[5-(2,6-dimethylphenyl)-3-(2,6-dimethylphenylethyl)pentyl]-1H-imidazole

4.0 g of 1-benzyl-5-[5-(2,6-dimethylphenyl)-3-hydroxy-3-(2,6-dimethylphenylethyl)pentyl]-1H-imidazole hydrochloride and 20 g of potassium hydrogen sulfate is combined and the mixture is heated for 6 hours at 150°C.

25 Ethanol (40 ml) is added and the mixture is filtered. 20 ml of conc. hydrochloric acid is added and the mixture is hydrogenated palladium on carbon (10 %) as catalyst until the hydrogen consumption ceases. The reaction mixture is filtered, water is added and the mixture is made alkaline with sodium hydroxide. The product is then extracted into toluene, which is washed with water, and evaporated to dryness. The residue, which is the product as base, is converted to nitrate with nitric acid in water. M.p. 147-150°C.

30

Example 10

4-[3,3-bis(3,5-dimethylphenyl)propyl]-1H-imidazole

35 a) 1-benzyl-5-[3,3-bis(3,5-dimethylphenyl)-3-hydroxypropyl]-1H-imidazole

1.06 g of magnesium turnings are covered with 30 ml of dry tetrahydrofuran. To the mixture is then added dropwise a solution of 5-bromo-m-xylene (8,14 g) in 10 ml of dry tetrahydrofuran at such a rate that a smooth reaction is maintained. After the addition is complete, the reaction mixture is refluxed for one additional hour and cooled to room temperature. The reaction mixture is then added dropwise to a solution of 1-benzyl-5-imidazole propionic acid ethyl ester (5,0 g) in 40 ml of tetrahydrofuran at 60°C. After the addition is complete, the reaction mixture is refluxed for 2 hours, cooled and poured into cold water. Tetrahydrofuran is evaporated and to the solution is added conc. hydrochloric acid. The solution is cooled, some ether is added and the precipitate which contains the product as hydrochloride salt is removed by filtration, washed and dried. Yield 4,1 g. M.p. 120-124°C.

45

b) 1-benzyl-5-[3,3-bis(3,5-dimethylphenyl)propen-2-yl]-1H-imidazole

4.0 g of 1-benzyl-5-[3,3-bis(3,5-dimethylphenyl)-3-hydroxypropyl]-1H-imidazole is dissolved in 30 ml of ethanol and 2 ml of conc. hydrochloric acid is added. The reaction mixture is then refluxed for 4 hours and evaporated to dryness. The residue which is the product is recrystallized from ethyl acetate. Yield 3,1 g M.p. 170-176°C.

50 According to the same procedure as the example the following substituted derivatives were prepared:
1-benzyl-5-(3,3-diphenylpropen-2-yl)-1H-imidazole, hydrochloride M.p. 173-175°C.

1-benzyl-5-[3,3-bis(2-methoxyphenyl)propen-2-yl]-1H-imidazole, hydrochloride. M.p. 191-194°C.

1-benzyl-5-[3,3-bis(3-methoxyphenyl)propen-2-yl]-1H-imidazole, hydrochloride. M.p. 132-135°C.

55 1-benzyl-5-[3,3-bis(4-methoxyphenyl)propen-2-yl]-1H-imidazole, hydrochloride. M.p. 157-163°C.

1-benzyl-5-[3,3-bis(2,3-dimethylphenyl)propen-2-yl]-1H-imidazole, hydrochloride.

¹H NMR (as base):

2.055 (s, 3H), 2.159 (s, 3H), 2.251 (s, 6H), 3.467 (d, 2H), 4.781 (s, 1H), 5.281 (s, 2H), 5.761 (t, 1H), 6.8-7.4 (m, 12H), 9.97 (s, 1H)

60 1-benzyl-5-[3,3-bis(2-methylphenyl)propen-2-yl]-1H-imidazole, hydrochloride. m.p. 84-87°C.

1-benzyl-5-[3,3-bis(3-methylphenyl)propen-2-yl]-1H-imidazole, hydrochloride. M.p. 115-117°C.

c) 1-benzyl-5-[3,3-bis(3,5-dimethylphenyl)propyl]-1H-imidazole

1-benzyl-5-[3,3-bis(3,5-dimethylphenyl)propen-2-yl]-1H-imidazole hydrochloride is dissolved in ethanol and a catalytic amount of Pd/C (10 %) is added. The reaction mixture is agitated vigorously at room temperature in

a hydrogen atmosphere until the uptake of hydrogen ceases. The mixture is filtered and the filtrate is evaporated to dryness. The residue which is the product is purified by flash chromatography eluating with methylene chloride-methanol mixture.

By the same method is prepared for example 1-benzyl-5-[3,3-bis(3-methoxyphenyl)propyl]-1H-imidazole hydrochloride, m.p. 165-167°C, and 1-benzyl-5-[3,3-diphenylpropyl]-1H-imidazole hydrochloride, m.p. 160-162°C.

5

d) 4-[3,3-bis(3,5-dimethylphenyl)propyl]-1H-imidazole

2,0 g of 1-benzyl-5-[3,3-bis(3,5-dimethylphenyl)propyl]-1H-imidazole hydrochloride is hydrogenated in the mixture of 30 ml of 2 N hydrochloric acid and 10 ml ethanol at 80°C Pd/C (10 %) as catalyst. When the uptake of hydrogen ceases, the reaction mixture is cooled, filtered and evaporated to dryness. Water is added and the mixture is made alkaline with sodium hydroxide. The product is then extracted to ethylacetate which is washed with water, dried with sodium sulfate and evaporated to dryness. The residue is the product as base and it is made to its hydrochloride salt in ethyl acetate using dry hydrochloric acid. Yield 0,6 g. M.p. of the product is 101-105°C.

10

¹H NMR: 2.247 (s, 12H), 2.2-3.7 (m, 4H), 3.798 (t, 1H), 4.788 (s, 2H), 6.8-7.2 (m, 6H), 7.214 (s, 1H), 8.715 (s, 1H)

15

Using the same method for example the following compounds included in the invention were prepared: 4-[3,3-bis(2,3-dimethylphenyl)propyl]-1H-imidazole

10

¹H NMR (as base):

20

2.097 (s, 6H), 2.260 (s, 6H), 2.3 (m, 2H), 2.6 (m, 2H), 4.389 (s, 1H), 6.0 (s, 1H), 6.712 (s, 1H), 7.011 (s, 6H), 7.508 (s, 1H)

20

4-[3,3-bis(2-methoxyphenyl)propyl]-1H-imidazole, hydrochloride. M.p. 194-196°C.

4-[3,3-bis(3-methoxyphenyl)propyl]-1H-imidazole

¹H NMR (as base):

25

2.5 (m, 4H), 3.747 (s, 6H), 3.862 (t, 1H), 6.6-7.3 (m, 9H), 7.498 (s, 1H), 8.165 (s, 1H)

4-[5-(2,6-dimethylphenyl)-3-hydroxy-3-(2,6-dimethylphenylethyl)pentyl]-1H-imidazole, hydrochloride. M.p. 178-180°C.

25

4-[3,3-bis(4-methoxyphenyl)propyl]-1H-imidazole

25

¹H NMR (as base):

25

2.5 (m, 4H), 3.744 (s, 6H), 3.815 (t, 1H), 6.1 (broad signal, 1H), 6.732-7.171 (m, 9H), 7.489 (s, 1H)

25

4-[3,3-bis(4-methylphenyl)propyl]-1H-imidazole

¹H NMR (as hydrochloride):

30

2.260 (s, 6H), 2.5 (m, 4H), 3.879 (t, 1H), 4.907 (s, 2H), 6.9-7.2 (m, 9H), 8.727 (s, 1H)

Example 11

35

1-benzyl-5-[3,3-bis(4-methylphenyl)propen-2-yl]-1H-imidazole

To a dry flask is placed 4,8 g (0,2 mol) of NaH (washed free from oil with cyclohexane). Onto it is then dropped 100 ml of dry dimethylsulfoxide. The reaction vessel is warmed at 80°C until the evolution of hydrogen ceases. The resulting solution of methylsulfinyl carbanion is cooled in an ice-water bath and 54,1 g of 3-(1-benzyl-5-imidazolyl)-propyltriphenylphosphonium bromide is added in 200 ml of dimethylsulfoxide. The reaction mixture is then stirred at room temperature for 0,5 hours and to it is added in small portions 23,0 g of 4,4'-dimethylbenzophenone. The reaction mixture is stirred at room temperature for 1 hour and some of the dimethylsulfoxide is distilled. The residue is poured into water which is made alkaline with sodium hydroxide. The product is extracted into toluene which is washed with water, dried with sodium sulfate and evaporated to dryness. From the residue which contains the crude product as base is converted into the hydrochloride in ethylacetate. Yield 32 g. M.p. 216-220°C.

40

¹H NMR: 2.289 (s, 3H), 2.370 (s, 3H), 3.467 (d, 2H), 4.764 (s, 1H), 5.302 (s, 2H), 6.030 (t, 1H), 6.8-7.4 (m, 9H), 8.9 (s, 1H)

45

50

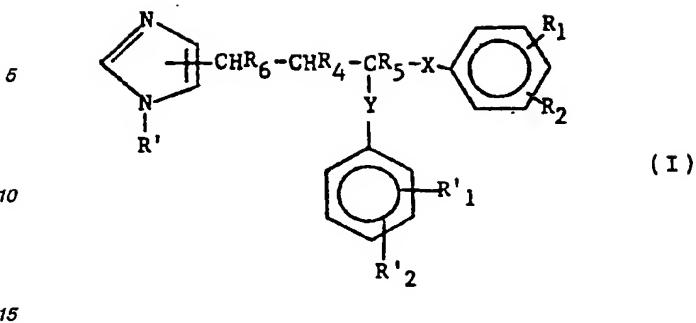
Claims

55

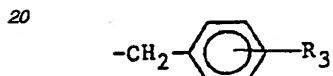
1. A substituted imidazole of the formula

60

65



or a non-toxic pharmaceutically acceptable acid addition salt thereof wherein R₁, R₂, R'₁ and R'₂, which can be the same or different, are H, CH₃, C₂H₅, OCH₃, OH, CH₂OH, NH₂ or halogen; R' is H or

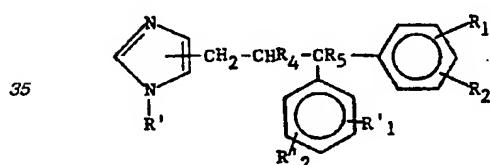


25

where R₃ is H, CH₃ or halogen; R₄ is H, R₅ is H or OH and R₆ is H or OH or one of R₅ and R₆ is H and the other, together with R₄, forms a bond and X and Y, which can be the same or different, are a bond, a straight C₁₋₂-alkyl or the corresponding alkenyl.

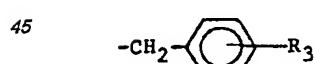
2. A substituted imidazole of the formula

30



40

or a non-toxic pharmaceutically acceptable acid addition salt thereof wherein R₁, R₂, and R'₁ and R'₂ which can be the same or different are H, CH₃, C₂H₅, OCH₃, OH, CH₂OH, NH₂ or halogen; R' is H or



50

where R₃ is H, CH₃ or halogen; R₄ is H and R₅ is H or R₄ and R₅ together form a bond.

3. A substituted imidazole according to claim 2 wherein R₄ and R₅ are both H.

4. A substituted imidazole according to claim 2 wherein R₄ and R₅ together form a bond.

5. A substituted imidazole according to claim 2 3 or 4 wherein at least one of R₁, R₂, R'₁ and R'₂ is not H and one or more of the substituents R₁, R₂, R'₁ and R'₂ are in the 3, 5, 3' or 5' positions of the phenyl groups.

6. A substituted imidazole according to claim 5 wherein R₂ and R'₂ both are H and R₁ and R'₁ are both not H and are both in the meta position of the phenyl groups.

7. A substituted imidazole according to any one of claims 2 to 6 wherein R' is H.

8. A substituted imidazole according to any one of claims 2 to 6 wherein R' is

60

65



5

where R3 is H, CH₃ or halogen.

9. A substituted imidazole according to claim 8 wherein R3 is H.

10. A compound according to claim 1 which is 4-[5-(2,6-dimethylphenyl)-3-hydroxy-3-(2,6-dimethylphenylethyl)pentyl]-1H-imidazole, 10
4-[3,3-bis(4-chlorophenyl)-3-hydroxypropyl]-1H-imidazole,
4-[3,3-diphenyl-3-hydroxypropyl]-1H-imidazole,
4-(3,3-diphenylpropen-2-yl)-1H-imidazole,
4-(3,3-diphenylpropyl)-1H-imidazole, 15
4-[3,3-bis(2-methylphenyl)-3-hydroxypropyl]-1H-imidazole,
4-[3,3-bis(4-chlorophenyl)propen-2-yl]-1H-imidazole,
4-[3,3-bis(2-methylphenyl)propen-2-yl]-1H-imidazole,
4-[3,3-bis(2-methylphenyl)propyl]-1H-imidazole,
1-benzyl-4-(3,3-diphenylpropyl)-1H-imidazole,
1-benzyl-5-(3,3-diphenylpropyl)-1H-imidazole, 20
4-[3,3-bis(3-methylphenyl)propyl]-1H-imidazole,
4-[3,3-bis(3-methylphenyl)propen-2-yl]-1H-imidazole,
1-(4-chlorobenzyl)-4-(3,3-diphenylpropyl)-1H-imidazole,
1-(4-chlorobenzyl)-5-(3,3-diphenylpropyl)-1H-imidazole,
4-[5-(2,6-dimethylphenyl)-3-(2,6-dimethylphenylethyl)pentyl]-1H-imidazole, 25
4-[3,3-bis(3,5-dimethylphenyl)propyl]-1H-imidazole,
4-[3,3-bis(3-methoxyphenyl)propyl]-1H-imidazole,
4-[3,3-bis(2,3-dimethylphenyl)propyl]-1H-imidazole,
1-benzyl-5-[3,3-bis(3-methoxyphenyl)propen-2-yl]-1H-imidazole, 30
1-benzyl-5-[3,3-bis(3,5-dimethylphenyl)propyl]-1H-imidazole,
4-[3,3-bis(4-methylphenyl)propyl]-1H-imidazole,
4-[3,3-bis(3-fluorophenyl)propyl]-1H-imidazole, 35
1-benzyl-5-[3,3-bis(4-chlorophenyl)-3-hydroxypropyl]-1H-imidazole,
1-benzyl-5-[3,3-bis(4-chlorophenyl)propen-2-yl]-1H-imidazole,
4-[3-(4-chlorophenyl)3-hydroxy-3-phenylpropyl]-1H-imidazole,
4-[3,3-bis(3-methylphenyl)-3-hydroxypropyl]-1H-imidazole,
4-[3,3-bis(3-fluorophenyl)propen-2-yl]-1H-imidazole, 40
1-benzyl-5-[5-(2,6-dimethylphenyl)-3-hydroxy-3-(2,6-dimethylphenylethyl)pentyl]-1H-imidazole,
1-benzyl-5-[3,3-bis(3,5-dimethylphenyl)-3-hydroxypropyl]-1H-imidazole,
1-benzyl-5-[3,3-bis(3,5-dimethylphenyl)propen-2-yl]-1H-imidazole, 45
1-benzyl-5-[3,3-bis(2-methoxyphenyl)propen-2-yl]-1H-imidazole,
1-benzyl-5-[3,3-bis(4-methoxyphenyl)propen-2-yl]-1H-imidazole,
1-benzyl-5-[3,3-bis(2,3-dimethylphenyl)propen-2-yl]-1H-imidazole,
1-benzyl-5-[3,3-bis(2-methylphenyl)propen-2-yl]-1H-imidazole, 50
1-benzyl-5-[3,3-bis(3-methylphenyl)propen-2-yl]-1H-imidazole,
4-[3,3-bis(2-methoxyphenyl)propyl]-1H-imidazole,
4-[3,3-bis(4-methoxyphenyl)propyl]-1H-imidazole,
1-benzyl-5-[3,3-bis(4-methylphenyl)propen-2-yl]-1H-imidazole,
or 1-benzyl-5-(3,3-diphenylpropen-2-yl)-1H-imidazole or a non-toxic pharmaceutically acceptable acid
addition salt thereof.

11. A pharmaceutical composition comprising a substituted imidazole as claimed in any one of claims 1 to 10 and a pharmaceutically acceptable carrier.

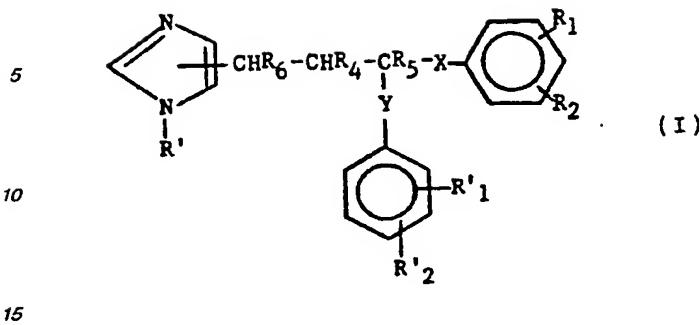
12. An imidazole derivative as claimed in any one of claims 1 to 11 or a non-toxic acid addition salt for use in therapy as an aromatase inhibiting agent. 55

Claims for the following Contracting States: ES, GR

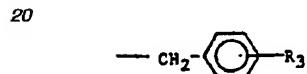
1. A process for the preparation of a substituted imidazole of the formula

60

65

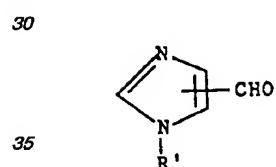


or a non-toxic pharmaceutically acceptable acid addition salt thereof wherein R₁, R₂, R'₁ and R'₂, which can be the same or different, are H, CH₃ C₂H₅, OCH₃, OH, CH₂OH, NH₂ or halogen; R' is H or



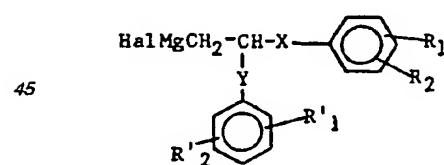
25

where R₃ is H, CH₃ or halogen; R₄ is H, R₅ is H and R₆ is H or OH or R₄ and R₆ together form a bond and X and Y which can be the same or different, are a bond, a straight C₁₋₂-alkyl or the corresponding alkenyl, which comprises reacting an 4(5)-imidazole aldehyde of the formula

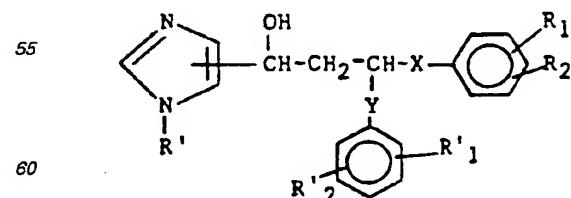


40

with a Grignard reagent



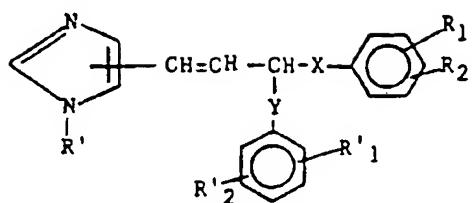
to give a compound of formula



65

wherein R', R₁, R₂, R'₁, R'₂, X and Y are as defined before; dehydrating this product to give a compound

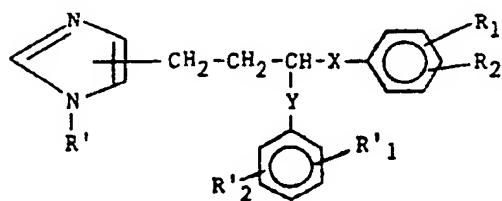
of formula



5

10

wherein R', R₁, R₂, R'₁, R'₂, X and Y are as defined before; and hydrogenating the product to give a compound of formula

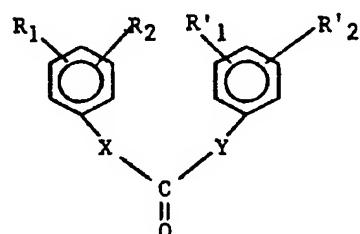


20

25

wherein R', R₁, R₂, R'₁, R'₂, X and Y are as defined before.

2. A process for the preparation of a compound of formula (I) as defined in claim 1 or a non-toxic pharmaceutically acceptable acid addition salt thereof wherein R₆ and R₄ are each H and R₅ is OH, which comprises reacting a ketone of the formula

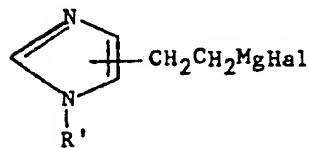


30

40

45

with a Grignard reagent



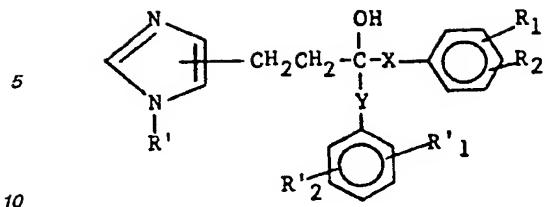
50

55

to give a compound of formula (I) which is

60

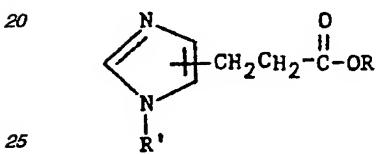
65



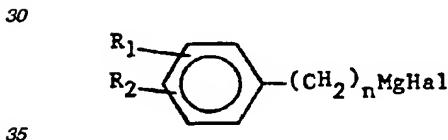
wherein R', R₁, R₂, R'1, R'2, X and Y are as defined before.

15

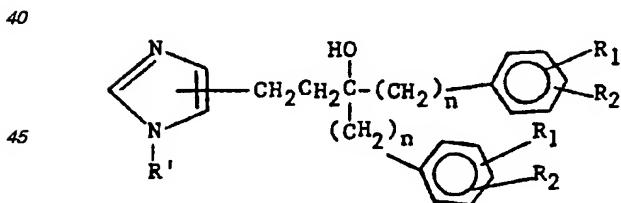
3. A process for the preparation of a compound of formula (I) as defined in claim 1 in which R₆ and R₄ are H, R₅ is OH and R₁ and R'1 are identical and R₂ and R'2 are identical or a pharmaceutically acceptable acid addition salt thereof, which comprises reacting an 4(5)-imidazole propionic acid alkyl ester of the formula



wherein R' is as defined in claim 1 and R is alkyl with a Grignard reagent of the formula

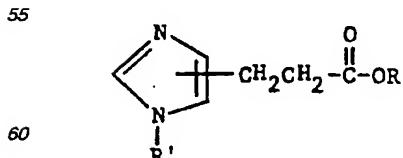


where R₁ and R₂ are as defined in claim 1 and n is 0 to 2 to give a compound of formula (I) which is



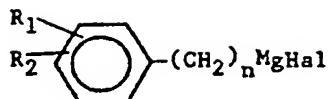
50

4. A process for the preparation of a compound of formula (I) as defined in claim 1 in which R₆ and R₄ are H and R₅ is OH or a non toxic pharmaceutically acceptable acid addition salt thereof which comprises reacting an imidazole ester of the formula



65

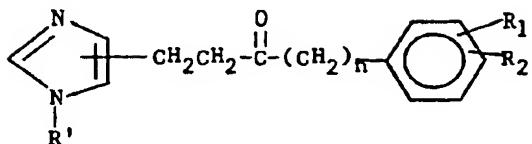
with a Grignard reagent



5

to give a compound of the formula

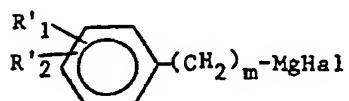
10



15

which further is reacted with another Grignard reagent of the formula

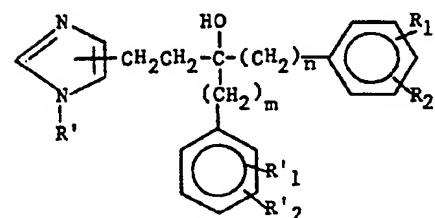
20



25

where R'1 and R'2 are as defined before and m is 0 to 2 to give a compound of formula (I) which is

30



35

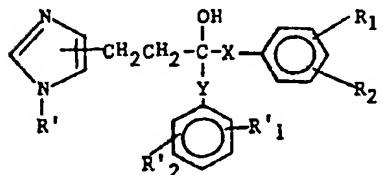
40

5. A process according to claim 3 or 4 wherein an 4(5)-imidazole propionic acid amide is used as starting material instead of the 4(5)-imidazole propionic acid ester.

6. A process for the preparation of a compound of formula (I) as defined in claim 1 in which R6 is H and R4 and R5 together form a single bond or a non-toxic pharmaceutically acceptable acid addition salt thereof

50

which comprises dehydrating a compound of formula

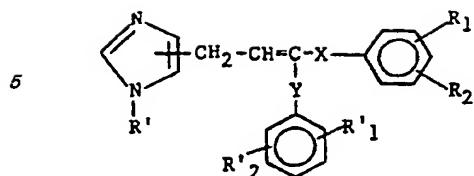


55

60

to give a compound of formula (I) which is

65

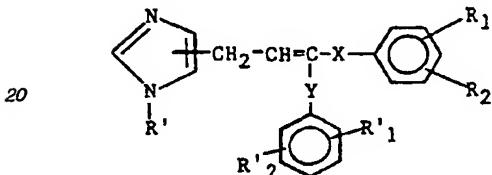


10

wherein R', R₁, R₂, R'₁, R'₂, X and Y are as defined before.

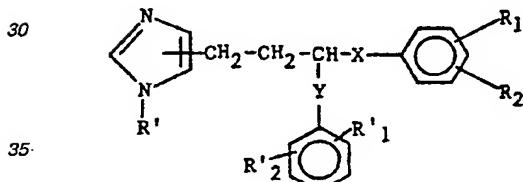
7. A process for the preparation of a compound of formula (I) as defined in claim 1 in which R₄, R₅, R₆ are each H or a non-toxic pharmaceutically acceptable acid addition salt thereof which comprises hydrogenating a compound of formula

15



25

to give a compound of formula (I) which is

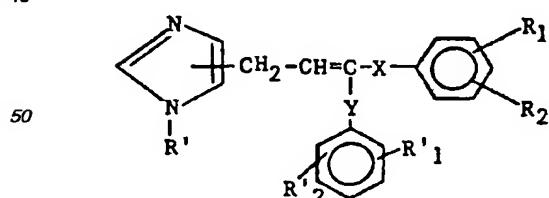


40

wherein R', R₁, R₂, R'₁, R'₂, X and Y are as defined before.

8. A process for the preparation of a compound of formula (I) as defined in claim 1 in which R₄, R₅, R₆ and R' are each H or a non-toxic pharmaceutically acceptable acid addition salt thereof which comprises hydrogenating a compound of formula

45

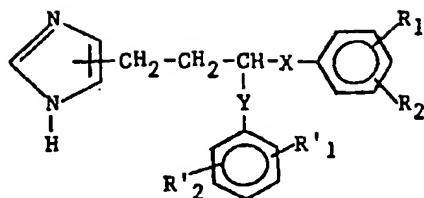


55

to give a compound of formula (I) which is

60

65



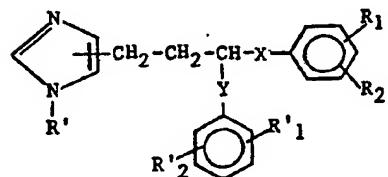
5

10

wherein R₁, R₂, R'₁, R'₂, X and Y are as defined before.

9. A process for the preparation of a compound of formula (I) as defined claim 1 in which R₄, R₅, R₆, and R' are each H or a non-toxic pharmaceutically acceptable acid addition salt thereof
which comprises hydrogenating a compound of formula

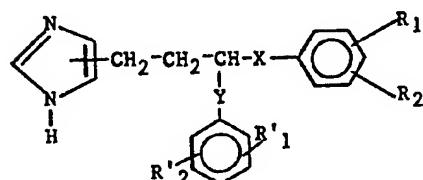
15



20

25

to give a compound of formula (I) which is



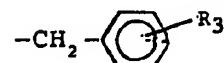
30

35

wherein R₁, R₂, R'₁, R'₂, X and Y are as defined before.

10. A process for the preparation of a compound of formula (I) as defined in claim 1 in which R' is

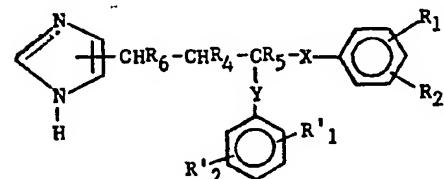
40



45

in which R₃ is as defined in claim 1 or a non-toxic pharmaceutically acceptable acid addition salt thereof
which comprises reacting a compound of the formula

50

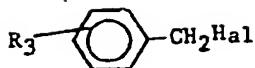


55

60

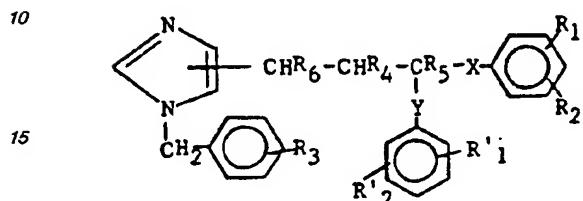
wherein R₁, R₂, R'₁, R'₂, R₄, R₅, R₆, X and Y are as defined before with a strong base and further with a
benzyl halide of the formula

65



5

where R₃ is as defined before to give a compound of formula (I) which is

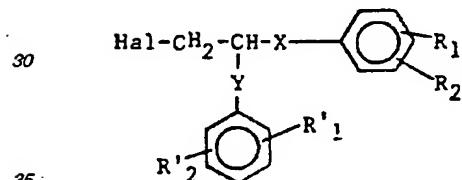


20

11. A process for the preparation of a compound of formula (I) as defined in claim 1 in which R₅ is H and R₄ and R₆ together form a single bond or a non-toxic pharmaceutically acceptable acid addition salt thereof

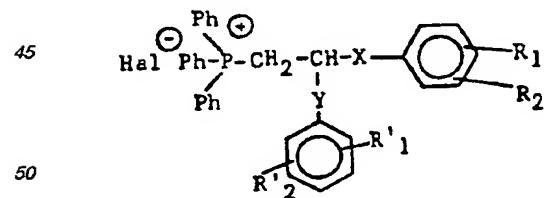
25

which comprises reacting a halogenated hydrocarbon of the formula



40

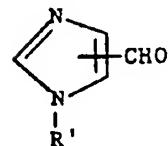
wherein R₁, R₂, R'₁, R'₂, X and Y are as defined before with triphenylphosphine to give a phosphonium salt of the formula



55

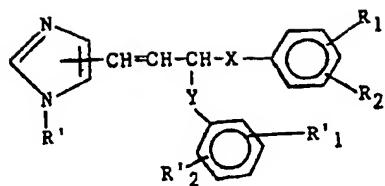
which is further reacted with a strong base and then with an 4(5)-Imidazole aldehyde of the formula

60

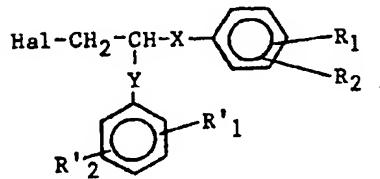


65

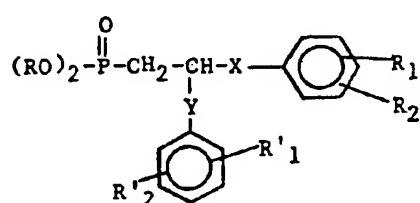
where R' is as defined before to give a compound of formula (I) which is



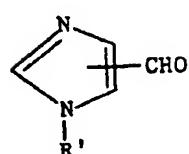
12. A process for the preparation of a compound of formula (I) as defined claim 1 in which R₅ is H and R₄ and R₈ together form a single bond or a non-toxic pharmaceutically acceptable acid addition salt thereof which comprises reacting a halogenated hydrocarbon of the formula



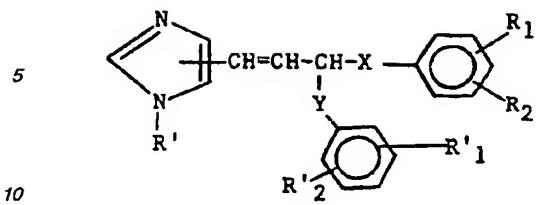
wherein R₁, R₂, R'1, R'2, X and Y are as defined before with a triester of phosphonic acid to give a compound of formula



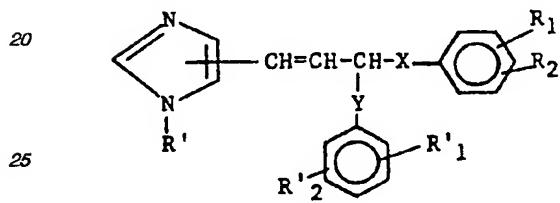
wherein R is alkyl, which further is reacted with a base and then with an 4(5)-imidazole aldehyde of the formula



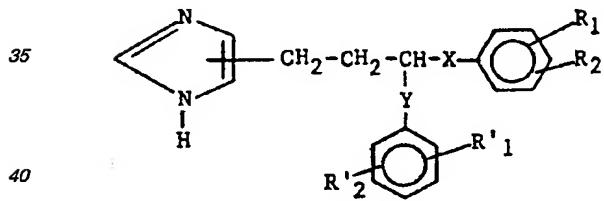
where R' is as defined before to give a compound of formula (I) which is



13. A process for the preparation of a compound of formula (I) as defined in claim 1 in which R₄, R₅, R₆, and R' are each H or a non-toxic pharmaceutically acceptable acid addition salt thereof which comprises
15 hydrogenating a compound of formula



30 to give a compound of the formula (I) which is



45 wherein R₁, R₂, R'1, R'2, X and Y are as defined before.

50

55

60

65



European Patent
Office

EUROPEAN SEARCH REPORT

Application Number

EP 88 30 9413

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. Cl. 4)
A	EP-A-0 194 984 (CONTINENTAL PHARMA) ---		C 07 D 233/64 C 07 D 233/58 A 61 K 31/415
A	EP-A-0 165 779 (ELI LILLY) ---		
A	EP-A-0 072 615 (FARMOS GROUP) ---		
A	EP-A-0 064 820 (FARMOS-YHTYMA OY) ---		
A	EP-A-0 058 047 (FARMOS-YHTYMA OY) ---		
A	EP-A-0 034 474 (FARMOS-YHTYMA OY) ---		
A	EP-A-0 034 473 (FARMOS-YHTYMA OY) -----		
			TECHNICAL FIELDS SEARCHED (Int. Cl.4)
			C 07 D 233/00 A 61 K 31/00
<p>The present search report has been drawn up for all claims</p>			
Place of search	Date of completion of the search	Examiner	
THE HAGUE	23-12-1988	DE BUYSER I.A.F.	
CATEGORY OF CITED DOCUMENTS		<p>T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document</p>	
<p>X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document</p>			